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(71) Applicant *(for all designated States except US)*: ALZA CORPORATION [US/US]; 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US).

(71) Applicant *(for US only)*: ECKENHOFF, Bonnie, J. (legal representative of the deceased inventor) [US/US]; 1080 Autumn Lane, Los Altos, CA 94022 (US).

(72) Inventor: ECKENHOFF, James, B. (deceased).

(72) Inventors; and

(75) Inventors/Applicants *(for US only)*: HOLLADAY, Leslie, A. [US/US]; Apartment 96, 1200 Dale Avenue, Mountain View, CA 94040 (US). LEONARD, John, Joseph, Jr. [US/US]; 11236 La Jolla Court, Cupertino, CA 95014 (US). LEUNG, Iris, K., M. [GB/US]; 1629 Bowling Lane, San Jose, CA 95118 (US). TAO, Sally, A. [US/US]; 1151 Miller Avenue, San Jose, CA 95129 (US). MAGRUDER, Judy, A. [US/US]; 355 Fay Way, Mountain View, CA 94043 (US). CARR, John, P. [US/US]; 376 Waverly Street, Sunnyvale, CA 94086 (US). WRIGHT, Jeremy [US/US]; 631 Cuestra Drive, Los Altos, CA 94024 (US).

(74) Agents: DILLAHUNTY, Mary, Ann et al.; Alza Corporation, 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US).

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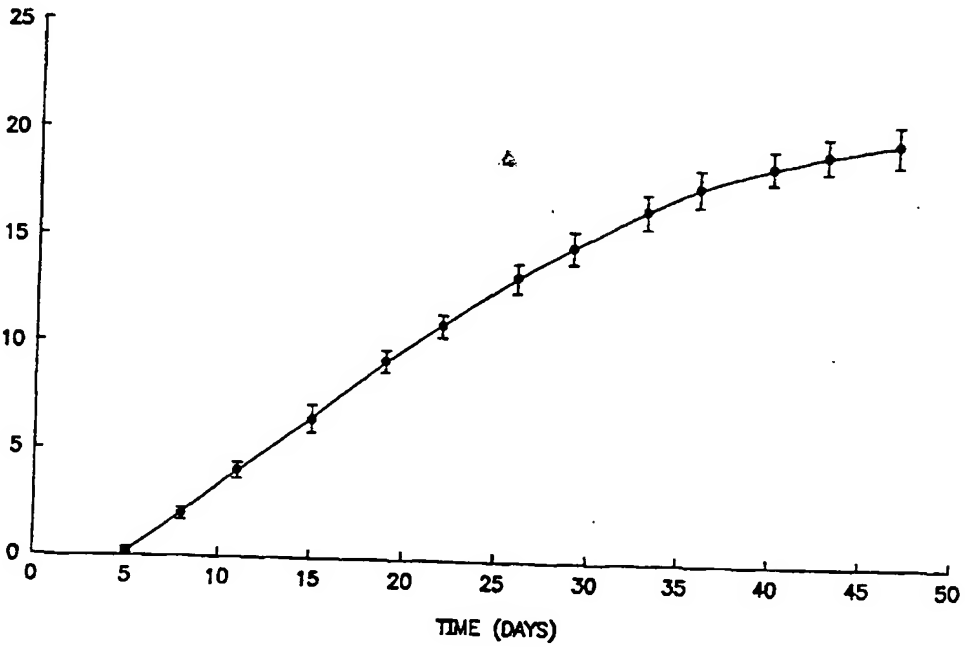
(54) Title: PEPTIDE/PROTEIN SUSPENDED FORMULATIONS



(57) Abstract

The present invention provides improved compositions for improving the chemical and physical stability of peptides and proteins. The invention provides a liquid beneficial agent formulation containing a liquid suspension comprising at least 5 % by weight beneficial agent and having a viscosity and beneficial agent size which minimizes setting of the agent in suspension over the extended delivery period.

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PEPTIDE/PROTEIN SUSPENDING FORMULATIONS

TECHNICAL FIELD

This invention relates to stabilized, concentrated suspensions formulations of peptides and proteins. More particularly, this invention relates to novel and improved compositions for providing concentrated, non-aqueous suspensions of peptides/proteins for pharmaceutical use having adequate chemical, physical and bioactive stability suitable for long term delivery from a sustained release drug delivery system.

BACKGROUND ART

Proteins, as well as many other biologically active compounds, degrade over time in aqueous solution. Because of this chemical instability, protein solutions are often not suitable for use in drug delivery devices. Carriers, in which proteins do not dissolve but rather are suspended, can often offer improved chemical stability. Furthermore, it can be beneficial to suspend the beneficial agent in a carrier when the agent exhibits low solubility in the desired vehicle. However, suspensions can have poor physical stability due to settling and agglomeration of the suspended beneficial agent. The problems with non-aqueous carriers tend to be exacerbated as the concentration of the active compound is increased.

For drug delivering implants, dosing durations of up to one year are not unusual. Beneficial agents which have low therapeutic delivery rates are prime candidates for use in implants. When the device is implanted or stored, settling of the beneficial agent in the liquid formulation can occur. This heterogeneity can adversely effect the concentration of the beneficial agent dispensed. Compounding this problem is the size of the implanted beneficial agent reservoir. Implant reservoirs are generally on the order of

25-250 μl. With this volume restriction, a formulation of high concentration (greater than or equal to 10%) and a minimum amount of suspension vehicle and other excipients is preferred.

Alpha interferon (α-IFN) is one example of a beneficial agent which provides a therapeutic effect at a low dose. This interferon is indicated in the treatment of chronic hepatitis because of its antiviral activity. Prescribed therapy presently entails injections of α-IFN solution, containing about 3.0×10^6 IU (15 micrograms) of agent per dose, three times per week for a 4 to 6 month period. Frequent injections are required because of the short elimination half-life of α-IFN; most of the drug being completely cleared from the plasma within eight to ten hours after the injection.

U.S. Pat. Nos. 4,871,538 issued to Yim et al; 4,847,079 issued to Kwan et al; 5,081,156 issued to Yamashira et al, and European Publication No. 0,281,299 issued to Yim et al describe IFN /peptide compositions with concentrations between 10^4 to 10^8 IU/ml. In Kwan et al, a pharmaceutical solution having a α-IFN concentration of 10^3 to 10^8 IU/ml is described. Yim describes a dosage range being between 10^4 to 10^8 IU α-IFN/ml. In Yim II, an insoluble complex including α-IFN, zinc, and protamine is suspended in a phosphate buffer. Yim I, Yim II, and Kwan, however, teach the use, in part, of an aqueous buffer in their compositions. This leads to possible hydrolysis of the compound, leading to chemical degradation and instability. Yamashira teaches a sustained release preparation of interferon in a mixture with a biodegradable carrier. IFN is incorporated at concentrations of 10^3 to 10^8 IU per 1 mg of carrier or, alternatively, each dosage form containing 10^4 to 10^8 IU of interferon. Furthermore, while the patents and publications described above describe concentrations between 10^4 to 10^8 IU/ml, none describe concentrations on the order of 10^9 to 10^{11} IU/ml.

There is a need for a novel composition comprising a nonaqueous suspension vehicle and concentrated protein/peptide as the beneficial agent for use in implanted, sustained release devices. While it is known in the art to achieve stable α IFN concentrations of up to 10^8 IU/ml, this invention utilizes a novel combination whose combined effect produces a significant and surprising improvement in the physical and chemical stability of the beneficial agent compound over other formulations.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a cross-section of an implantable sustained release osmotic delivery device for use in combination with the concentrated suspensions of the present invention.

FIG. 2 is a graph illustrating the stability of a cytochrome c suspension.

FIG. 3 is a graph illustrating the stability of an α-interferon suspension.

DESCRIPTION OF THE INVENTION

One aspect of this invention relates to preparations for stabilizing peptides and proteins at high concentrations for extended periods of time.

Another aspect of this invention relates to stabilized preparations of human α-IFN.

Another aspect of this invention relates to stabilized preparations of human α-IFN having concentrations of at least 1×10^9 IU/ml.

Another aspect of this invention relates to stabilizing beneficial agent formulations comprising a beneficial agent having a particle size of between 0.3 to 50 microns and suspension vehicle formula having a viscosity between 100 to 100,000 poise at 37°C.

The new formulations are physically stable suspensions which provide chemical stability to water sensitive compounds and can be employed to stabilize high concentrations of the active compound. The carrier components are acceptable for use in implantable systems.

MODES FOR CARRYING OUT THE INVENTION

The concentrated beneficial agent suspensions of the present invention provide significantly stable concentrations over extended periods of time, useful for sustained delivery, implant applications. The suspensions of this invention minimize the particle degradation due to hydrolysis and particle settling over the duration of the extended delivery period. These extended periods of time are between one week to two years, preferably between three months to one year.

The sustained parenteral delivery of drugs provides many advantages. Typical sustained release implantable osmotic delivery devices are described in U.S. Pat. Nos. 5,034,229; 5,057,318; and 5,110,596 which are incorporated herein by reference. As shown in Fig. 1, these devices 10 typically comprise a housing 12 including a fluid impermeable wall section 14 and a fluid permeable wall section 6 which sections define and surround an internal compartment 18. An exit passageway 20 is formed within the fluid impermeable wall section to fluidly communicate the internal compartment 18 with the external environment. To minimize exposure to the environmental fluids, a beneficial agent 22 is contained within the fluid impermeable section. An expandable driving member 24, contained within the fluid permeable section, expands with the imbibition of fluid across the fluid permeable wall section. Typically a piston 26 separates the beneficial agent 22 from the expandable driving member 24. This forces the agent out through the exit

passageway and into the environment of use. The non-aqueous administration of a beneficial agent in the suspension formulation as disclosed herein can be accomplished using implant devices of these kinds.

According to this invention, high concentrations of the beneficial agent remain suspended, and physically and chemically stable in a non-aqueous suspension vehicle. "High concentration" is defined as the beneficial agent concentration level of at least about 0.5 wt% of the formulation, preferably at least about 5 wt% and most preferably between about 10 to 70% w/w. For example, "high concentrations" of α-IFN are 10^9 to 10^{11} IU; and for salmon calcitonin, concentrations of between 2×10^4 IU to 2.8×10^6 IU are "high concentrations". The beneficial agent particle size is between 0.3 to 50 microns, and preferably about 1-10 microns in diameter. Desired particle size can be provided typically by milling, sieving, spray drying, supercritical fluid extraction of the particular beneficial agent selected. Typical beneficial agents for use in this device and composition include the interferons and calcitonin. Other representative beneficial agents that can be administered include pharmacologically active peptides and proteins, anabolic hormones, growth promoting hormones, hormones related to the endocrine system comprising porcine growth promoting hormone, bovine growth promoting hormone, equine growth promoting hormone, ovine growth promoting hormone, human growth promoting hormone, growth promoting hormones derived by extraction and concentration from pituitary and hypothalmus glands, growth promoting hormones produced by recombinant DNA methods, bovine growth promoting hormone as described in Nucleic Acid Res., Vol. 10, p 7197 (1982), ovine growth promoting hormone as described in Arch. Biochem. Biophys., Vol. 156, p 493 (1973), and porcine growth promoting hormone as described in DNA, Vol. 2, pp 37, 45, (1983). Representative beneficial agents also comprise cochicine, cosyntropin, and lypressin. The polypeptides also comprise growth hormone, somatropin, somatotropin, somatotropin analogues, modified porcine somatotropin,

modified bovine somatotropin, derivatives of both porcine and bovine somatotropin, somatomedin-C, gonadotropic releasing hormone, follicle stimulating hormone, luteinizing hormone, LH-RH, LH-RH analogs, growth hormone releasing factor, gonadotropin releasing factor, insulin, chorionic gonadotropin, oxytocin, somatotropin plus an amino acid, vasopressin, adrenocorticotrophic hormone, epidermal growth factor, prolactin, somatostatin, somatotropin plus a protein, polypeptides such as thyrotropin releasing hormone, thyroid stimulating hormone, secretin, pancreozymin, enkephalin, glucagon, endocrine agents secreted internally and distributed in an animal by way of the bloodstream, and the like. The beneficial agents and their dosage unit amounts are known to the prior art in The Pharmacological Basis of Therapeutics, by Gilman, Goodman, Rall and Murad, 7th Ed., (1985) published by MacMillan Publishing Co., NY; in Pharmaceutical Sciences, Remington, 17th Ed., (1985) published by Mack Publishing Co., Easton, PA, and in U.S. Pat. No. 4,526,938. Particularly preferred are beneficial agents which produce the desired therapeutic effect at a low delivery rate/dose, for example, proteins/peptides which require picograms to milligrams of agent.

A pharmaceutically acceptable suspension vehicle is used to suspend the solid beneficial agent particles in the beneficial agent formulation. Non-aqueous vehicles are used to isolate the beneficial agent from water and prevent hydrolysis or other degradation of the beneficial agent while in suspension. Furthermore, pharmaceutically acceptable suspension vehicles may function as a thickening agent for the components present in an implant. As a vehicle for transporting beneficial agents from the implant, it provides protection against the decomposition of a beneficial agent, and it imparts physical and chemical stability to components present in the formulation. The thickening agent may be used to increase the viscosity of the formulation to prevent fluids in the implantation environment from mixing with the

implant's beneficial agent formulation. The amount of thickening agent present in the formulation is between 1% to 99.9% and preferably 5-60% depending upon the viscosity adjustment needed.

Typical non-aqueous suspension vehicles include: waxes, which have a softening temperature at or less than body temperature; hydrogenated vegetable oils, (e.g., peanut oil, cottonseed oil, sesame oil, castor oil, olive oil, corn oil, Iodinated poppy seed oils) silicon oil, medium chain fatty acid monoglycerides, or polyols. Of these polyols are preferred.

Polyols suitable for suspension vehicles include such as diol, triol, polyhydric alcohol, and the like. More specific polyols comprise polyethylene glycol (average molecular weight between 200 and 1000), propylene glycol, polyethylene glycol 1,5-pentylene glycol; 1,6-hexylene glycol; 1,7-heptylene glycol; 1,9-nonylene glycol; 1,2-dimethyl-1,6-hexylene glycol; 1,2,3-propanetriol; 1,2,5-pentanetriol; 1,3,5-pentanetriol; 1,2,4-butanetriol; dipentaerythriol, and the like. In another embodiment the pharmaceutically acceptable suspension vehicle comprises glycerol mono(lower alkyl) ethers and glycerol di(lower alkyl) ethers such as glycerol 1-methyl ether; glycerol 1-ethyl ether; glycerol 1,2-dimethyl ether; glycerol 1,3-dimethyl ether, and the like. In another embodiment the pharmaceutically acceptable vehicle comprises a mixture such as propylene glycol and glycero, and the like.

Sufficient viscosity is required to suspend the particles in the carrier throughout the duration of the extended delivery period. Settling is a function of the particle size and the carrier viscosity. If the duration of the delivery period is shorter, the viscosity can be lower since the time required to be

suspended is shorter. The viscosity required, for example, can be determined by the Stokes-Einstein equation which is a measure of how far a particle in suspension will travel

$$V = \frac{2gR^2 (P_p - P_c)}{9\mu}$$

V = velocity of settling
μ = viscosity of the carrier
g = acceleration due to gravity
P_p = density of particle
P_c = density of carrier

wherein R = the average particle radius of the beneficial agent. The viscosity of the beneficial agent suspending formulation can be altered by the use of thickening agents to raise the viscosity to the desired level. Typical thickening agents for use in the compositions of this invention include suitable hydrogels such as hydroxypropyl cellulose, hydroxypropyl methyl cellulose (HPMC), sodium carboxymethyl cellulose, polyacrylic acid, poly(methyl methacrylic acid) (PMMA). Preferred hydrogels are cellulose ethers such as hydroxyalkylcellulose and hydroxyalkylalkyl-cellulose compounds. A most preferred hydroxyalkylcellulose is hydroxypropyl cellulose (HPC) and povidone (PVP). Hydroxypropyl cellulose is commercially available in a wide range of viscosity grades sold under the tradename Klucel ™ (Hercules, Ltd., London, England). The concentration of the hydroxyalkylcellulose is dependent upon the particular viscosity grade used and the desired viscosity of the liquid composition. For example, where the desired viscosity is less than about 1000 poise (cps), hydroxypropyl cellulose having an average molecular weight of about 60,000 daltons (i.e., Klucel EF ™) can be used. Where the desired viscosity is from about 1000 to about 2500 cps, higher viscosity grades of hydroxypropyl cellulose can be used (i.e., Klucel LF ™ and Lucel GF ™). In addition to using different viscosities of different thickening

agents, using different amounts of the same particular thickening agent can be used to vary the viscosity. Preferably, the concentration of hydroxypropyl cellulose is from 5 percent w/w and, more preferably from 5 to 20 %w/w of the carrier and most preferably between 8-18 %w/w. Aluminum monostearate can be used as a thickening agent if oils are used as the carrier.

Hydroxyalkylalkylcellulose ethers are a class of water-soluble hydrogels derived from etherification of cellulose. As used herein in reference to this class of hydrogels, the term "alkyl" means C_1-C_6 alkyl where alkyl refers to linear or branched chains having 1 to 6 carbon atoms, which can be optionally substituted as herein defined. Representative alkyl groups include methyl, ethyl, propyl, isopropyl, butyl, pentyl, hexyl and the like.

Exemplary hydroxyalkylalkylcelluloses are hydroxypropylmethyl cellulose, hydroxyethylmethyl cellulose and hydroxybutylmethyl cellulose. Hydroxypropylmethyl cellulose (HPMC) is preferred. HPMC is commercially available (i.e., Aldrich Chem. Co., Ltd. Dorset, England and Dow Chem. Co., Midland, Mich., USA) in a wide range of viscosity grades. In addition to increasing viscosity, hydroxyalkylalkylcelluloses can serve as a stabilizing, suspending and emulsifying agent. The concentration of hydroxyalkylalkylcellulose in a liquid composition of this invention is dependent inter alia on its intended use (i.e., stabilizer, emulsifier, viscosity-increasing agent) and its viscosity grade.

To assure the viscosity of the suspension vehicle is sufficient to maintain the agent in suspension over the desired delivery period, thickening agents can be added to the suspension vehicle. The preferred thickening agents include povidone and hydroxypropyl cellulose. In one embodiment, when the PEG utilized is a low molecular weight, e.g., 400, 5% hydroxypropyl cellulose, having an average molecular weight of 1000, or 40 -60% povidone can be used in combination with a balance of polyethylene glycol. If the

polyethylene glycol utilized in the suspension vehicle has a molecular weight of greater than 600, e.g., 1000 molecular weight, povidone is preferably utilized as the thickening agent.

The following examples are offered to illustrate the practice of the present invention and are not intended to limit the invention in any manner.

EXAMPLE 1

A viscous carrier was prepared containing 50% PEG 400 and 50% povidone (PVP) by weight. PEG 400 (Union Carbide) was weighed into a beaker and an equal weight of povidone K29-32 (GAF) was added. The PEG and povidone were mixed by stirring with a spatula for about 5 minutes. The blended carrier was allowed to sit overnight to insure complete dissolution of the povidone. The carrier was then deaerated in a vacuum oven (National Appliance Company) by drawing a vacuum and holding the carrier at 50°C for 30 minutes.

Cytochrome c (Sigma, from horseheart) was milled in a jar mill and then passed through a 400 mesh screen to produce a particle diameter of less than 37 micron. In a beaker, 0.5566 grams of the cytochrome c was added to 4.9970 grams of the PEG 400/povidone carrier to prepare a 10% cytochrome c suspension in 50:50 PVP:PEG 400 carrier. The suspension was thoroughly blended by mixing with a spatula for about 5 minutes. The cytochrome c suspension was then loaded into 11 osmotic veterinary implants (as in Figure 1).

The implants were tested in vitro by releasing into culture tubes filled with deionized water. To monitor release of cytochrome c from the implants, samples of the release media were assayed on a UV spectrophotometer (Shimadzu UV 160U) at a wavelength of 409 nm. The implants delivered the cytochrome c successfully over the designed duration of the implant

(42 days). Fig. 2 is a graph that illustrates the cumulative protein delivery (mg) over time. During the later half of the release period, several implants were removed from the tubes and examined to determine whether settling of the cytochrome c had occurred. These implants were sectioned and samples of the protein suspension were removed from the top and bottom portions of the implant. The samples of the protein suspension were weighed, diluted with DI water in volumetric flasks and assayed via UV a spectrophotometer. Results indicated that the cytochrome c suspension was homogeneous.

EXAMPLE 2

Standard : 20 µl of a 8.0 mg/ml standard was diluted to 160 µg/ml. Each HPLC sample was diluted by a factor of 10 into distilled water. The operating conditions of the HPLC were as follows:

Column:	POROS RH 2.1 mm x 3.0 cm
Mobile phase:	A: 95% H2O, 0.1% TFA, 5% ACN
	B: 95% ACN, 5% H2O, 0.083% TFA
Gradient:	20% B to 50% B in 5 minutes
Flow:	2.0 ml/min
Detector:	280 nm @ 0.002 AUFS

IRMA Standards: Working standards were prepared by diluting IRMA standards into phosphate buffered saline (PBS) containing 0.5% Bovine Serum Albumin (BSA). Samples were prepared by serially diluting by factors of 400 for interferon formulations and 2000 for the standard into PBS containing 0.5% BSA.

Figure 3 shows the results of the HPLC and the IRMA assays. The HPLC measurements indicate no losses of the α-IFN over 5 days, even at 37° C, indicating stability of this protein in non-aqueous vehicle. Relative to the initial stock solution, the activity shown by IRMA at t = 0 is 78%.

At t = 5 days, the formulation displayed an activity of 87% at room temperature and 90% at 37°C. When compared to the original stock, no losses of α-IFN were detected by HPLC in this formulation. Stability of interferon in PEG over 5 days at 37° C was indicated by this assay. However, approximately 80 - 90% of the activity of the initial stock was maintained. The IRMA readings suggest no activity losses due to time and temperature effects.

This invention has been described in detail with particular reference to certain preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

WHAT IS CLAIMED IS:

1. A beneficial agent formulation for use in a device which delivers the formulation over an extended delivery period, the formulation comprising a suspension containing at least 5% by weight beneficial agent in the form of solid particles, the beneficial agent particle size being 0.3 to 50 microns and the suspension viscosity being sufficient to prevent settling of the agent in the suspension formulation over the extended delivery period.

2. The formulation of claim 1, wherein the particle size is between 1 to 10 microns.

3. The formulation of claim 1, wherein the viscosity is 100 to 100,000 poise at 37°C.

4. The formulation of claim 1, wherein the extended delivery period is at least about 1 month.

5. The formulation of claim 1, wherein the liquid suspension further comprises a low molecular weight polyol and a thickening agent.

6. The formulation of claim 5, wherein the polyol is polyethylene glycol having a molecular weight between 200 and 1000.

7. The formulation of claim 6, wherein the thickening agent comprises povidone.

8. The formulation of claim 5, wherein the polyol is polyethylene glycol having a molecular weight between 200 and 600.

9. The formulation of claim 8, wherein the thickening agent comprises povidone or hydroxypropyl cellulose.

10. The formulation of claim 1, wherein the beneficial agent is human α-interferon.

11. The formulation of claim 10, wherein the concentration of interferon is at least 1×10^9 IU.

12. The formulation of claim 1, wherein said beneficial agent is a water sensitive compound.

13. A beneficial agent delivery device containing the formulation of claim 1.

14. The beneficial agent delivery device of claim 13, wherein the device is adapted to be implanted within an animal.

15. A composition for sustained controlled delivery over an extended delivery period, the composition comprising:

(a) 0.5% to 70% by weight beneficial agent having a particle size of between 0.3 to 50 microns; and

(b) a non-aqueous liquid suspension formulation characterized by a viscosity of between 100 to 100,000 poise at 37 ° C, the formulation further comprising polyethylene glycol with a molecular weight between 200 and 1000 and a thickening agent.

16. The composition according to claim 15, wherein the thickening agent comprises povidone or hydroxypropyl cellulose.

17. A beneficial agent delivery device containing the composition of claim 15.

18. The beneficial agent delivery device of claim 17, wherein the device is adapted to be implanted within an animal.

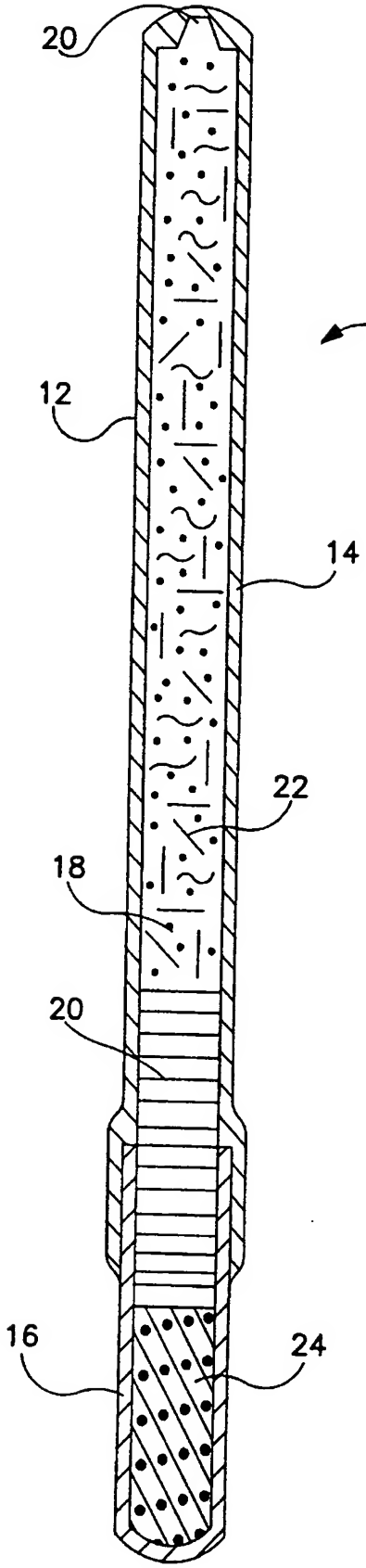
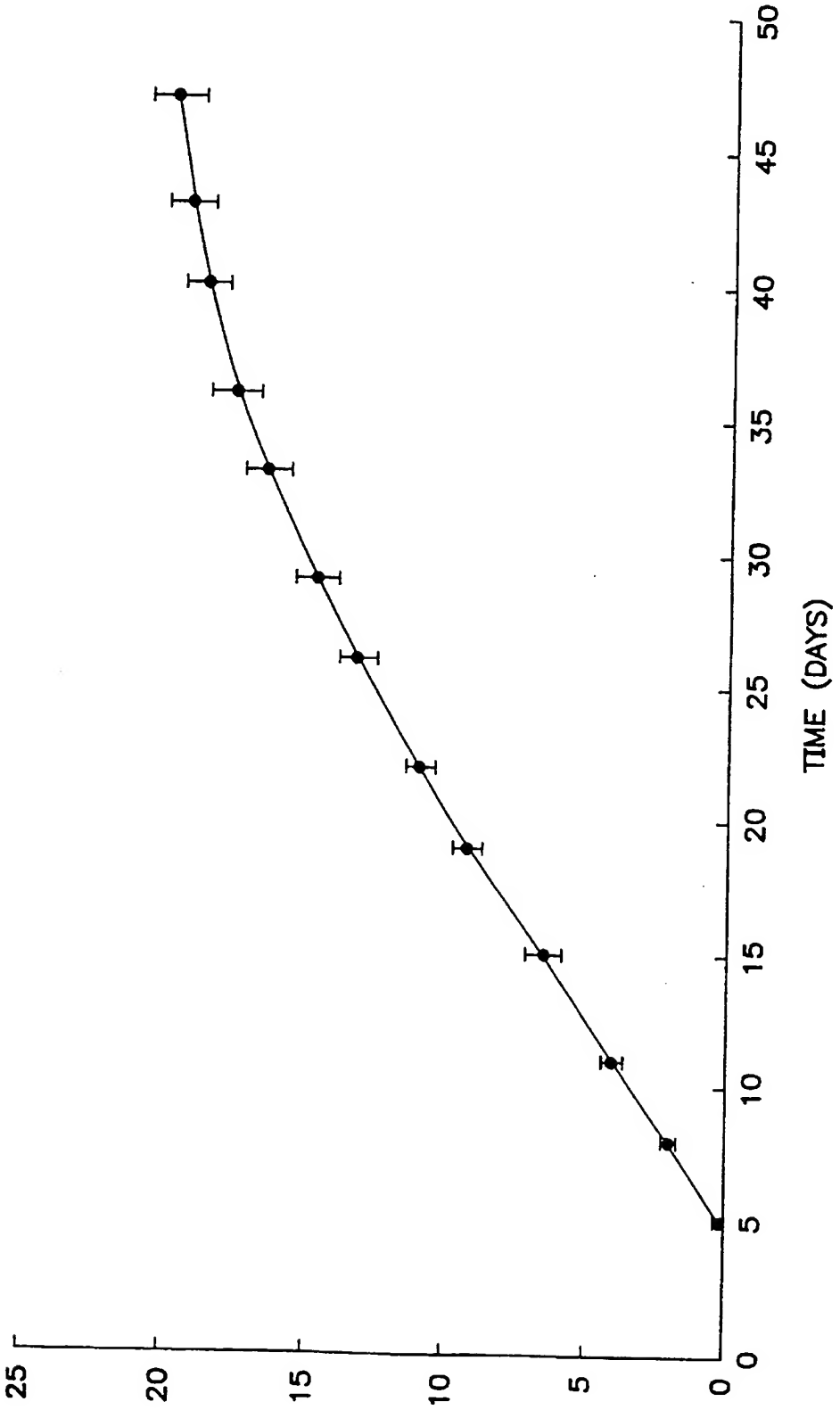


FIG. 1

FIG. 2



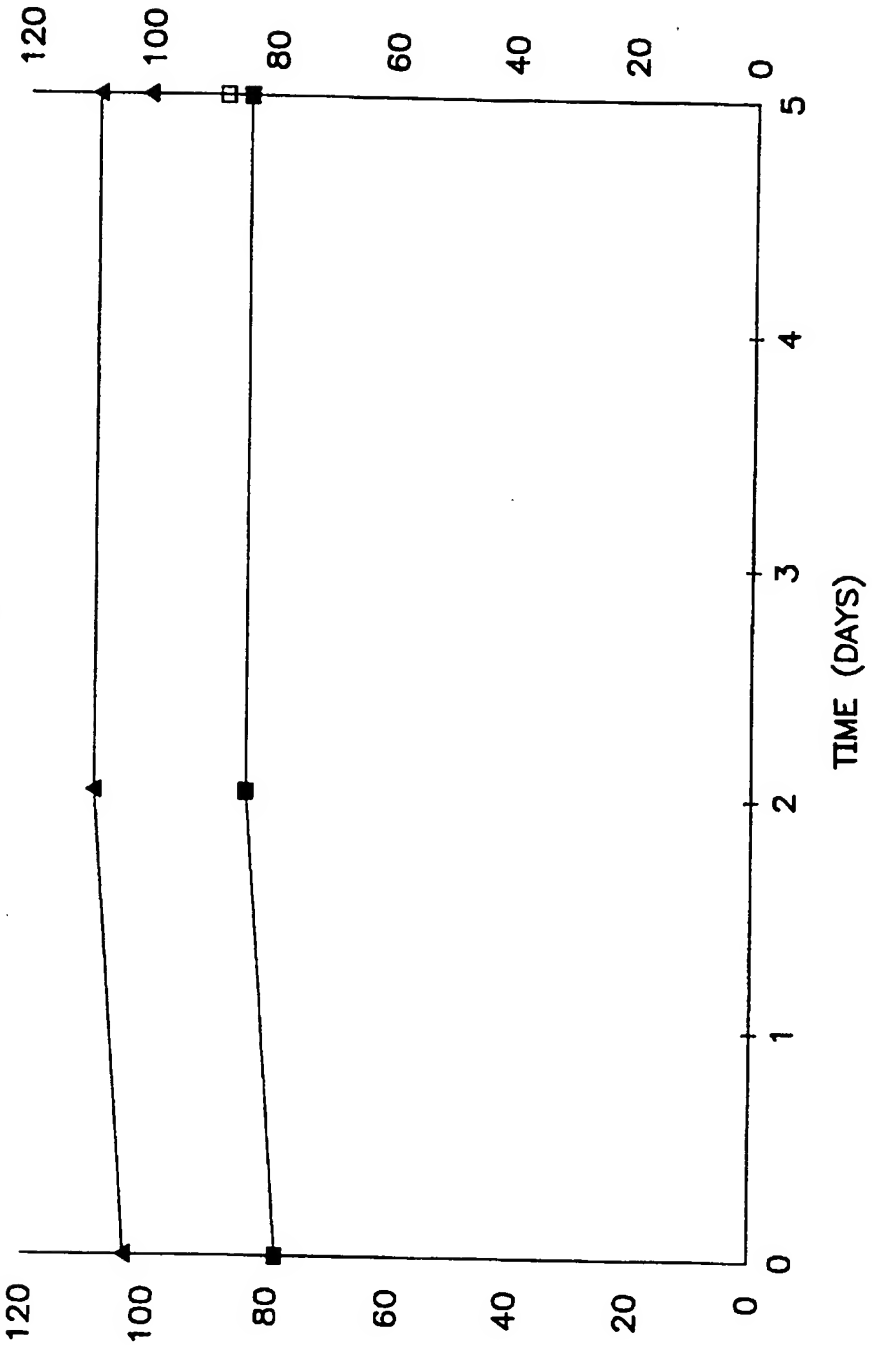


FIG. 3

INTERNATIONAL SEARCH REPORT

Inte .nal Application No
PCT/US 96/07377

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61K9/00 A61K47/10 A61K47/32 A61K47/38

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X Y	EP,A,0 374 120 (MONSANTO COMPANY) 20 June 1990 see claims 1,7,8 see page 6, line 54 - page 7, line 3 see page 7, column 19 - column 21 see page 7, column 38 - column 40 see page 7, column 50 - column 52 ---	1,3,4, 12-14 10
X	US,A,4 855 141 (ALZA CORPORATION) 8 August 1989 see claim 1 see column 10, line 22 - column 11, line 15 --- -/--	1,5,12, 13

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

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Date of the actual completion of the international search	Date of mailing of the international search report
17 September 1996	30.09.96

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INTERNATIONAL SEARCH REPORT

Int. ional Application No
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,Y	WO,A,95 34285 (ALZA CORPORATION) 21 December 1995 see claims 1,2 see page 14, line 20 - page 16, line 2 see page 20, line 3 - line 11 -----	10

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INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte. nal Application No
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Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP-A-374120	20-06-90	AU-A-	4617689	21-06-90
		CA-A-	2005226	13-06-90
		CZ-A-	8907028	18-01-95
		JP-A-	2204418	14-08-90
US-A-4855141	08-08-89	AR-A-	240398	30-04-90
		AU-A-	3012189	28-09-89
		CA-A-	1337038	19-09-95
		DE-T-	68907769	04-11-93
		EP-A-	0337613	18-10-89
		IE-B-	61335	02-11-94
		JP-A-	1299568	04-12-89
		NO-B-	177887	04-09-95
		PT-B-	90074	31-03-94
		US-A-	4959218	25-09-90
		US-A-	4996060	26-02-91
WO-A-9534285	21-12-95	AU-A-	7108094	05-01-96